



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁵ : C12N 9/42, 15/56 // C12N 9/42 C12R 1:885	A1	(11) International Publication Number: WO 93/24622 (43) International Publication Date: 9 December 1993 (09.12.93)
(21) International Application Number: PCT/FI93/00219 (22) International Filing Date: 24 May 1993 (24.05.93) (30) Priority data: 922373 22 May 1992 (22.05.92) FI 931193 17 March 1993 (17.03.93) FI (71) Applicant (for all designated States except US): VALTION TEKNIILLINEN TUTKIMUSKESKUS [FI/FI]; P.O. Box 402, FIN-02151 Espoo (FI). (72) Inventors; and (75) Inventors/Applicants (for US only) : BUCHERT, Johanna [FI/FI]; Westendintie 93 I 55, FIN-02160 Espoo (FI). SIIKA-AHO, Matti [FI/FI]; Susitie 26 E 33, FIN-00800 Helsinki (FI). VIKARI, Liisa [FI/FI]; Lökkikuja 5 F, FIN-00200 Helsinki (FI). PENTTILÄ, Merja [FI/FI]; Vähätuvantie 9 A 6, FIN-00390 Helsinki (FI). SALO- HEIMO, Anu [FI/FI]; Kyntäjätie 18 B, FIN-00390 Helsinki (FI). RANUA, Marijatta [FI/FI]; Kaarnaistenk- atu 30 A, FIN-02100 Lohja (FI).		(74) Agent: SEPPO LAINE KY; Lönnrotinkatu 19 A, SF- 00120 Helsinki (FI). (81) Designated States: AT, AU, BB, BG, BR, CA, CH, CZ, DE, DK, ES, FI, GB, HU, JP, KP, KR, LK, LU, MC, MN, MW, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SK, UA, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI pa- tent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>In English translation (filed in Finnish).</i>
(54) Title: MANNANASE ENZYMES, GENES CODING FOR THEM, METHODS FOR ISOLATING THE GENES, AND METHODS FOR BLEACHING LIGNOCELLULOSIC PULPS		
(57) Abstract The present invention relates to a DNA sequence, which codes for endomannanase produced by fungi of the genus <i>Tricho-</i> <i>derma</i> and transferred into a yeast or fungus strain induces that strain to produce endomannanase, as well as to a method for iso- lating genes coding for endomannanases. The invention also relates to vectors, yeast strains and fungal strains containing the DNA sequence. Furthermore, the invention provides an enzyme product containing at least one endomannanase, which contains at least one of the following endomannanases produced by fungi of the <i>Trichoderma</i> genus: an enzyme having mannanase activi- ty and an isoelectric point (pI) of about 3.8, an enzyme having mannanase activity and a pI of about 4.1, an enzyme having man- nanase activity and a pI of about 4.5, an enzyme having mannanase activity and a pI of about 5.4 and an enzyme having manna- nase activity and a pI of about 6.5, the isoelectric points being determined by isoelectric focusing. The endomannanase-enzyme and enzyme products according to the invention can be used for hydrolyzation of mannopolymers, in particular in connection with bleaching of lignocellulosic pulps.		

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Document Listing

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Mannanase enzymes, genes coding for them, methods for isolating the genes, and methods for bleaching lignocellulosic pulps.

5 The present invention relates to a DNA sequence which codes for mannanase enzymes. The invention also concerns vectors, yeast and fungus strains containing said DNA sequence. Furthermore the invention relates to a method for isolating genes coding for mannases and to a method for constructing yeast strains capable of expressing mannanase. The invention also provides an enzyme product containing at least one
10 mannanase and a method for preparing mannanase enzymes and enzyme products. Finally, the invention concerns a process for hydrolyzing mannopolymers and to a process for bleaching lignocellulosic pulps.

The main components of wood are cellulose, lignin and hemicellulose. Softwood
15 mainly comprises arabino-4-O-methylglucuronoxylan, whereas hardwood xylan consists of O-acetyl-4-O-methylglucuronoxylan. The glucomannan polymers are formed by a main chain consisting of glucose and mannose units. The main chain is substituted with galactose and acetylunits. There are only small amounts (2 - 5 %) of gluco-
mannanes in hardwood and the glucomannan of hardwood trees differs structurally
20 from that of softwood trees in the sense that it is not acetylated nor does it contain galactose side groups (Timell 1967). During chemical pulping the relative amounts of hemicelluloses is changed compared to that of the native tree. The main hemicellulose component of both softwood and hardwood kraft pulp is xylan (Sjöström 1977).
Softwood pulp contains also large amounts of glucomannan. During pulping, a part of
25 the hemicellulose of the wood is dissolved because of the very alkaline cooking liquor. When the cooking is continued, pH decreases and xylan with no or few side chains will start to precipitate on the surface of the cellulose fibers (Yllner and Enström 1956,
Yllner et al. 1957). This xylan reprecipitation takes place simultaneously with the precipitation of dissolved lignin. The glucomannan behaves in alkaline cooking
30 conditions in a different way compared to xylan. Softwood glucomannanes are subjected to a partial decomposition already at approx. 130 °C. and, thus, the relative proportion of glucomannan in pulp is smaller than in the original softwood (Sjöström

1977). The glucomannan remaining in the fiber is, however, very stable (Simonson 1971). Glucomannan is believed to be evenly distributed within the fiber, whereas the xylan concentration is largest on the surface of the fiber (Luce 1964).

- 5 After chemical pulping, the pulp will contain darkened residual lignin which is removed by bleaching. In conventional chlorine bleaching the lignin is dissolved using chlorine or chlorine dioxide. Nowadays oxygen bleaching, hydrogen peroxide and combinations of these and the former are also often used.
- 10 It has been found out that the bleachability of the fibres can be improved by using hemicellulases, in particular xylanase, (Kantelinen et al 1988, Viikari et al 1991a). The amounts of enzymes needed for achieving bleaching are small and an enzymatic treatment can easily be combined with pulping processes. In the present applications, thus far enhanced bleachability of the fibres has been achieved by hydrolyzing the fibre
- 15 xylan. According to the present belief, the xylanases act mainly on the xylan located on the surface of the pulp fibres (Kantelinen et al 1991). Hemicellulase (xylanase) treatments have been combined with various bleaching sequences. The use of enzymes in bleaching has been evaluated by peroxide delignification (Viikari et al. 1986, Viikari et al. 1987, Viikari et al. 1990) and by various chlorine bleaching sequences carried
- 20 out on laboratory scale. Industrially enzyme treatments have been combined with different chlorine bleaching sequences and also with various multistage peroxide bleaching sequences (Viikari et al. 1991b). However, even if the peroxide bleaching is carried out as a multistage bleaching sequence, the brightness of pulp bleached entirely without chlorine chemicals remains lower than that of chlorine bleached pulp.
- 25 The use of bacterial and fungal mannanases for pretreatment of pulp before bleaching has been studied (Clark et al 1990, Clark et al 1991). The mannanase preparations used have been produced by the organisms *Bacillus subtilis*, *T. harzianum* and *Aspergillus niger*. The products are reported slightly to improve bleaching but, because of the
- 30 impurities which the products contain, the influence of the mannanase component has not been shown.

It is an aim of the present invention to eliminate the problems associated with the prior art and to provide a completely novel enzyme product which can be used, for instance, in bleaching of cellulose pulp.

5 The present invention is based on the surprising finding that the mannanase enzymes produced by fungi of the genus *Trichoderma* are particularly well suited for hydrolyzing mannopolymers and, thus, for treatment of cellulose pulp, for instance.

10 It is known in the art that the mannanase production of several fungi can be induced by using cellulose (Lyr 1963) and mannan-containing components (Reese and Shibata 1964) in the growth medium. It is also known that the fungus *Trichoderma reesei* is an efficient producer of enzymes which hydrolyze cellulose (for instance Bisaria and Ghose 1981) and xylan (Poutanen et al. 1987). *Trichoderma reesei* growing in nature can utilize not only the cellulose of plant material but also the hemicellulose part.

15 In the research work forming the basis of the invention it could be shown that a certain strain of *Trichoderma reesei* (Rut C-30, VTT D-86271) produces a lot of mannanase activity when cultivated in a fermentor on a culture medium containing cellulose as a carbon source and corn steep syrup as a nitrogen source. In connection with the
20 invention, an enzyme mixture was obtained from the *Trichoderma reesei* strain which contained at least one selected from the group comprising an enzyme having mannanase activity and an isoelectric point (pI) of about 3.8, an enzyme having mannanase activity and a pI of about 4.1, an enzyme having mannanase activity and a pI of about 4.5, an enzyme having mannanase activity and a pI of about 5.4 and an enzyme having
25 mannanase activity and a pI of about 6.5. The isoelectric points indicated have been determined by using isoelectric focusing, the accuracy of the determination being on the order of $\pm 0,5$ pI units.

30 The properties of the above mannanase preparation differ from those of the known mannases and it is far more efficient in the hydrolysis of mannopolymers than, for instance, the mannanase isolated from the species *Bacillus subtilis*.

According to the invention the desired enzyme(s) can also be produced by other *Trichoderma reesei* strains, by other strains belonging to the genus *Trichoderma* and by using other culture media. For this reason the genes coding for the above-mentioned enzymes have, in connection with the present invention, been isolated in order to provide the possibility to produce the enzyme by using, for instance, genetically or mutationally improved *Trichoderma reesei* strains or by using other genetically modified production hosts (such as yeast cells) to which the genes coding for the *Trichoderma reesei* mannanases have been transferred.

A novel method for isolating the genes has, at the same time, been developed.

The characteristic features of the invention are indicated in the attached claims.

The term "mannanase" denotes an enzyme which is capable of cleaving polyose chains containing mannose units (mannopolymers). Mannanase therefore covers both endomannanases which internally cleave mannopolymers and exomannanases which cleave mannopolymers from the terminal ends of the chain. As examples of mannopolymers, glucamannan, galactoglucomannan and galactomannan may be mentioned.

Within the scope of the present application, the term "enzyme preparation" designates any product which contains at least one mannanase enzyme. Thus, the enzyme preparation can, for instance, be a culture medium containing a mannanase or several mannanases, an isolated mannanase, or a mixture of two or more mannanases.

By "hybridization" are meant conditions under which different forms of DNA sequences hybridize to DNA sequences coding for a *Trichoderma* mannanase.

According to the present invention, the mannanase was isolated from the culture medium of *Trichoderma reesei* by using known methods for purifying proteins. For isolating the enzymes anionic and cationic ionexchangers were used as was hydrophobic interaction based chromatography, the isolation of the enzyme being surprisingly easy

and fast. The invention is not, however, restricted to this enzyme isolation method, but it is possible to isolate the enzyme with other known methods.

5 The mannanase activity of the enzyme has been determined by using a method in which an enzyme sample is allowed to act on the mannan of Locust bean for a desired period of time, after which the sugars released are analysed. The method is the following:

10 A 0.5 % solution of Locust bean gum mannan (Sigma G-0753) is prepared in a 50 mM sodium citrate buffer at pH 5.3, and the insoluble solid matter is removed from this mannan solution by centrifugation. To 1.80 ml of the mannan solution 0.200 ml of a suitably diluted enzyme sample is added and it is allowed to act on mannan at +50 °C. Exactly 5 minutes after the addition of the enzyme sample 3.00 ml of DNS reagent is added and the mixture formed is heated for 5 min in boiling water. Then, the mixture is cooled with cold water. The colour formed is measured by using spectrophotometer
15 at 540 nm. The amount of reducing sugars in the sample is determined by subtracting from the value obtained by spectrophotometer the influence of the reducing sugars contained in the enzyme sample as such, and then comparing the value with values obtained when a mannose solution of known concentration is treated in the same way as the sample. One activity unit (cat/ml) corresponds to the amount of enzyme contained
20 in 1 ml of undiluted sample, which releases 1 mole of reducing sugars per second under the experimental conditions described above.

The invention further describes the specific genes encoding *Trichoderma* mannanase. Typically, the DNA sequence according to the invention codes for the mannanase of a
25 fungus of the genus *Trichoderma* and transferred to a yeast or fungal strain induces that strain to produce mannanase.

It should be noted that present knowledge on biochemistry and and molecular biology shows that an enzyme activity suitable for similar application can be provided also by
30 expressing parts of the above-mentioned genes or by expressing gene forms or synthetic genes which differ from the native genes as far as their nucleotide sequences are concerned. Therefore the scope of the invention also covers all forms of genes which

remind of the genes described in this invention and which code for a similar kind of mannanase activity.

5 Vectors, such as the yeast vectors pMAN1, pMAN2, pMAN3 or pMAN4 described in Example 1, can be formed from the DNA sequence. These vectors inserted into the yeast strain *Saccharomyces cerevisiae* DBY746 have been deposited with the collection Deutsche Sammlung von Mikroorganismen und Zellkulturen GmbH on 30 December 1992 under the numbers DSM 7363, DSM 7364, DSM 7366 and DSM 7365, respectively. The yeast strain used for cloning is described in more detail in the article
10 Penttilä, M.E. et al., Cloning of *Aspergillus niger* genes in yeast. Expression of the gene coding *Aspergillus* β -glucosidase, Mol. Gen. Genet. 194 (1984), 494-499.

The present invention also provides a method for isolating genes coding for mannanases of, in particular, *Trichoderma reesei*. The method is, however, suited for the isolation
15 of genes coding for mannanases of other microorganisms, as well. The method utilizes an expression library constructed in yeast, in particular in *Saccharomyces cerevisiae*, the genes of the fungus being expressed in the yeast under a yeast promoter. The mannanase protein is extracellularly secreted by the yeast and the yeast clones containing the mannanase gene can be found on basis of their production of mannanase activity, which can be detected by enzyme activity tests or by plate assays.
20

Alternatively, the yeast producing mannanase activity can be found, for instance, by using the antibody corresponding to the mannanase. The enzyme activity based method is a particularly preferred alternative for isolating mannanase genes because it makes it possible to find the yeasts producing the active mannanase enzyme and the genes can be
25 isolated from these yeasts for characterization. The yeast expression library can be constructed using some other yeast promoter, even a promoter providing a lower expression level. It is possible also to use the mannanase gene's own promoter, and the genes can be isolated by using a chromosomal gene library. The gene library can be constructed also, for instance, in a single-copy plasmid.

30

Based on the above, the method according to the invention for isolating genes coding for mannanases can be defined by the following preferred method steps:

The messenger RNA pool of a microorganism producing mannanase activity is enriched in respect of the messenger RNA of the mannanase by culturing the microorganism in conditions which will induce the mannanase production of said microorganism. The mannanase production is induced by, e.g., mannan-containing and/or cellulose-
5 containing culture media. Messenger RNA is isolated from the microorganism and the cDNA corresponding to the isolated mRNA is prepared. Then the cDNA thus obtained is placed in a vector under the control of a yeast promoter and the recombinant plasmids obtained are transformed into a yeast strain, which naturally does not produce the corresponding mannanase, in order to provide an expression library. The yeast
10 clones thus obtained are then cultured on a culture medium in order to express the expression library in the yeast. The yeast clones producing the mannanase are separated from the other yeast clones, and the plasmid-DNAs of said separated yeast clones are isolated. If desired, the DNA is sequenced in order to determine the DNA sequence coding for the mannanase.

15 As mentioned above, the mannanase-producing microorganism used in the method preferably comprises a fungus and in particular a fungal strain belonging to the genus *Trichoderma*. According to one preferred embodiment the fungus is cultivated on a culture medium comprising at least mannan- (e.g. galactomannan) and/or cellulose-
20 containing substrate as carbon source and in addition possibly acetyl glucurono xylan and/or xylan. As Example 2 below will show, the culture medium can for instance contain a combination of all these substances and as nitrogen source, for instance, corn steep syrup.

25 In the following step of the method the recombinant plasmids are preferably transferred to a yeast strain belonging to the genus *Saccharomyces*, in particular to the strain *Saccharomyces cerevisiae*, which is cultivated for producing mannanase. The yeast promoter of the plasmid vector can be, for instance, PGK or a similar promoter which is considered to be strong. The yeast clones can be cultivated on different substrates
30 such as glucose and galactose. The culture medium is selected depending on under which promoter the gene is placed. According to the invention, it is also possible to use a yeast vector, wherein the promoter can be induced by galactose, the hydrolase genes

being expressed by galactose and not by, for instance, glucose. The benefit of the promoter PGK is a strong level of expression which is advantageous as far as enzymes having a low catalytic activity are concerned. It is also easy to manipulate it. As described in more detail in Example 1, the plasmid vector used, can comprise, for instance, vector pAJ401, which is prepared from vector pFL60 by cutting it with restriction enzymes and ligating to it an adapter consisting of oligonucleotides. The cDNA synthesized from the messenger RNA is then ligated between the promoter and terminator of vector pAJ401, downstream from the promoter.

It should be noticed that, within the scope of the present invention, the mannanase genes can, in principle, be isolated by all other commonly known methods, for instance by constructing an expression library into an *E. coli* strain bacterium, e.g. into a lambda vector, whereby the clone containing the gene corresponding to a certain protein can be found by, e.g. the antibody corresponding to the protein. The gene can also be found by using oligonucleotides corresponding to the N-terminal amino acid sequence of the mannanase protein as a probe in gene library hybridization.

The present invention relates to the expression of mannanase genes of the fungus *T. reesei* in yeast and to the secretion of the active gene product into the culture medium of the yeast. Because *S. cerevisiae* normally does not produce mannanase activity, a recombinant yeast which produces *T. reesei* mannanase can be used as such for producing an enzyme composition containing a specific mannanase. The prior art is completely silent about the expression of mannanase genes in foreign host organisms. Our invention shows that fungal mannanase can be produced in yeast, which means that the invention includes the production of mannanase enzyme in other yeast strains as well, for instance in yeast strains belonging to the genera *Kluyveromyces*, *Pichia* and *Hansenula*. The mannanase gene can be transferred to other fungi, such as strains of the genera *Aspergillus*, *Penicillium*, *Neurospora* and *Phanerochaete*, by methods known per se and the fungus can be made to secrete active *Trichoderma* mannanase. Even the mannanase production of *Trichoderma* itself can be improved or modified after gene isolation by known gene technical means, by, for instance, transferring several copies of the chromosomal mannanase gene into the fungus or by placing the gene under the

(e.g. stronger) promoter of another gene and thus to provide mannanase expression under desired growth conditions, such as on culture media which natively do not produce mannanases. The invention covers the organisms producing said *Trichoderma* mannanases.

5

The recombinant yeast or fungal strain according to the invention is therefore characterized in that it contains a DNA sequence which codes for a mannanase of a fungus of the genus *Trichoderma*. The yeast strain is preferably a strain of the species *Saccharomyces cerevisiae*, such as S.c.man1, S.c.man2, S.c.man3 or S.c.man4 (cf. Example 1 below). The recombinant fungal strain is preferably a strain belonging to the genus *Trichoderma*.

10

Based on the above, the invention also provides a method for constructing yeast strains which are capable of expressing mannanase. According to the method DNA sequences coding for mannanase are isolated from a suitable fungus species, a yeast vector is formed from the DNA sequence and the yeast vector is transferred to a suitable yeast strain. For instance, the yeast vectors are transferred to a yeast strain selected from the group comprising strains of genera *Saccharomyces*, *Kluyveromyces*, *Pichia* or *Hansenula*.

15

20

The invention also comprises a method for producing mannanase enzyme, which method comprises isolating the DNA sequence coding for mannanase from a fungus of the strain *Trichoderma reesei*, forming a vector containing said DNA sequence, which is then transferred to a suitable yeast or fungus strain for achieving a recombinant strain. Said recombinant strain is then cultivated under conditions which enable the strain to express mannanase. The mannanse produced is recovered, for instance, by isolating it from the culture medium.

25

The mannanase preparations produced according to the invention can be advantageously used for hydrolysis of mannopolymers, in particular in connection with cellulosic pulp bleaching. The mannanses used in bleaching comprise mannanases isolated from *T. reesei* (or produced in any of the above-mentioned ways) either in the form of a

30

mixture, in purified form or in culture medium. The mannanases can be used together with xylanase. According to a particularly preferred embodiment a combined mannanase/xylanase treatment is carried out in connection with bleaching employing chlorine chemicals. Chlorine free bleaching processes (e.g. peroxide bleaching) provide rather good results already by a mannanase enzyme preparation. The enzyme treatment makes it possible even further to increase the extraction of residual lignin in chemical bleaching, and the method is therefore as well environmentally as economically advantageous.

By means of the mannanase treatment described in this invention it is possible to achieve an increase of the brightness produced by, in particular, totally chlorine free pulp bleaching sequences, which cannot be obtained with peroxide only or with any other chemical at so low costs or without impairing the strength of the pulp. The mannanase treatment according to the invention combined with a xylanase treatment improves brightness particularly when it is used in connection with both conventional chlorine-based bleaching sequences and totally chlorine chemical free bleaching sequences. In conventional chlorine bleaching, the pulp can be bleached with a standard dosage of the chemical to a higher final brightness or, alternatively, by adding enzymes it is possible to lower the chemical dosage in the initial stage of the bleaching when a certain brightness level is aimed at, which makes it possible to reduce the detrimental impact on the environment.

The higher efficiency of the enzymes make it possible directly to affect the type and amount of the chemicals used for industrially extracting lignin from the fibres and, thus, to improve the low-chlorine or chlorine-free bleaching methods which are environmentally advantageous.

The invention is described in more detail with the help of the following non-limiting examples.

In the Examples reference is made to the attached drawings, whereby Figure 1 shows the structure of plasmid pAJ401 constructed from plasmid pFL60,

Figure 2 shows the restriction sites of the mannanase-coding insert of yeast vector pMAN1,

Figure 3 shows the restriction sites of the mannanase-coding insert of yeast vector pMAN2,

5 Figure 4 shows the restriction sites of the mannanase-coding insert of yeast vector pMAN3, and

Figure 5 shows the restriction sites of the mannanase-coding insert of yeast vector pMAN4.

10 Example 1

Isolation of mannanase genes and expression in yeast

Trichoderma reesei -strain QM 9414 was cultivated in a fermentor for 42 h on a cultivation medium containing (per litre): 20 g of Solca floc cellulose, 10 g of Locust
15 bean gum galactomannan (Sigma), 5 g of KH_2PO_4 , and 5 g of $(\text{NH}_4)_2\text{SO}_4$. After the 42 hours of growth, 1 g of lactose, 1 g of Birke 150 acetylglucuronoxylan and 1 g of Oat Spelt xylan was added for each litre of cultivation medium, and the fungus was cultivated for further 24 h. The RNA was isolated from the fungus as described by Chirgwin *et al.* (1979), and the poly A+ mRNA was isolated by chromatography
20 through oligo(dT) essentially as described (Maniatis *et al.*, 1982). cDNA was synthesized by using the ZAP-cDNA synthesis kit from Stratagene and it was ligated to the plasmid vector pAJ401. Plasmid pAJ401 was derived from plasmid pFL60 (Minet & Lacroute, 1990) by cutting it with EcoRI and XhoI restriction enzymes and by ligating inbetween an adapter, which was obtained by combining the oligonukleotides
25 5'-tcgaagaattcgagagactcgagt-3' and 3'-tcttaagctctctgagctcattaa-5'. The orientation of the restriction sites of enzymes EcoRI and XhoI was then reversed in the plasmid. The synthesized cDNA was ligated to the pAJ401 vector cut with EcoRI and XhoI between the PGK promoter and terminator. The 5' end of the cDNA is bound downstream from the PGK promoter during cloning. The ligating mixture was transformed into the *E.*
30 *coli* strain PLK-F' by electroporation (Dower *et al.*, 1988).

The structure of plasmid pAJ401 is shown in the figure.

The gene library plasmids were isolated as one batch from about 42,000 bacterial colonies and transformed into the *Saccharomyces cerevisiae* yeast strain DBY746 (Penttilä *et al.*, 1984) by electroporation (Becker & Guarante 1990) while selecting the transformants on a Sc medium which is deficient in uracil. The yeast colonies were
5 scraped from the plates and maintained as one batch in a Sc-ura-medium containing 15 % glycerol at -70 °C.

For isolating the mannanase genes the gene library mixture was pipetted onto substrate-containing plates (yeast minimal medium plates, containing for each 1000 ml: 0.058 g
10 histidine, 0.082 g tryptophan, 0.262 g leucine and 0.1 % Locust bean gum (0.5 % stock solution in 50 mM citrate buffer at pH 5.3. In all about 100,000 gene library clones were tested for mannanase activity after one week of cultivation by replicating first the colonies from the plates, then flushing the plates with water, and by staining the plates with a 0.1 % solution of Congo Red. Yeast colonies producing hydrolysis
15 halos were picked up from the replica plates and they were tested again for mannanase activity on plate assay.

Four mannanase-positive yeast clones were called *Saccharomyces cerevisiae* man1, S.c. man2, S.c. man3 and S.c. man4. These strains were deposited into the collection
20 Deutsche Sammlung von Mikroorganismen und Zellkulturen GmbH on the 30th of December 1992 under the numbers DSM 7363, DSM 7364, DSM 7366 and DSM 7365, respectively.

The yeast clone S.c. man1 (DSM 7363) contains the yeast vector pMAN1, which
25 consists of the plasmid pAJ401, having a 0.7 kb mannanase coding insert in the restriction sites of restriction enzymes EcoRI and XhoI.

The yeast clone S.c. man2 (DSM 7364) contains the yeast vector pMAN2, which
30 consists of the plasmid pAJ401, having a 0.7 kb mannanase coding insert in the restriction sites of restriction enzymes EcoRI and XhoI.

The yeast clone S.c. man3 (DSM 7366) contains the yeast vector pMAN3, which

consists of the plasmid pAJ401, having a 1.4 kb mannanase coding insert in the restriction sites of restriction enzymes EcoRI and XhoI.

5 The yeast clone *S.c. man4* (DSM 7365) contains the yeast vector pMAN2, which consists of the plasmid pAJ401, having a 1.0 kb mannanase coding insert in the restriction sites of restriction enzymes EcoRI and XhoI.

10 The DNA of the mannanase positive yeast clones was isolated (Sherman et al 1981), and used for transformation of the *E. coli* strain DH5 α . From the ampicillin-positive transformants plasmid DNA was isolated by conventional methods and digested by several restriction enzymes, to determine which clones contained the same mannanase gene. The clones were sequenced by conventional methods to obtain a suitably long a clone from each gene.

15 The cDNA sequence of the mannanase genes was determined by conventional methods. The nucleotide sequence of the insert of the vector pMAN1 is indicated in sequence list SEQ NO. 1.

20 The chromosomal copies were also isolated from the clones using the previously prepared gene library (Vanhanen et al. 1989), and the 5' end of the gene was sequenced by utilizing primers specific for the 5' end designed on basis of the cDNA sequence. Sequence listing SEQ ID NO. 2 shows the nucleotide sequence of the 5' end and sequence listing SEQ ID NO. 3 the nucleotide sequence of its 3' end. Sequence listing SEQ ID NO. 4 indicates the 5' end of the insert of vector pMAN3 and sequence listing SEQ ID NO. 5 the 3' end thereof.

25

Example 2

Isolation of mannanase genes

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This example illustrates an alternative way of isolating mannanase genes from a gene library mixture. *Trichoderma reesei* strain QM 9414 was cultivated, the gene library

plasmids were formed and the genes were expressed in yeast as described in Example 1.

5 The gene library mixture was pipated into the Sc-ura cultivation medium on microtiter plates so that each well of the microtiter plates contains an average of 70 different yeast library clones, in total about 75,000 clones. The microtiter plates were cultivated for 4 days and the cells were centrifuged to the bottom of the wells. The supernatants were tested for their mannanase activity by using Locust bean gum as a substrate. (cd. above). Some of the wells contained mannanase activity, and the yeast cells contained
10 in said wells were plated onto Sc-ura-plates as separate colonies. About 130 separate colonies of each well were retested for their mannanase activity in order to identify single yeast clones containing the mannanase gene.

15 Example 3

Cultivation and characterization of the yeast clones

This example describes an alternative embodiment for characterization of yeast clones containing mannanase genes.

20

Yeast clones S.c.man1, S.c.man2, S.c.man3 and S.c.man4, which correspond to plasmid clones pMAN1, pMAN2, pMAN3 and pMAN4, were obtained by following the procedure described in Example 1. The clones were cultivated on a Sc-ura cultivation medium for 2 days in a flask and the supernatant was analysed by the
25 Western blot -method using antibodies prepared against purified mannanases obtained from *T.reesei*. The 5' end of the nucleotide sequence obtained from the clones was also compared with the N-terminal sequence from the purified mannanases. It was thus ascertained which isolated mannanase genes correspond to the earlier purified mannanase proteins.

30

Example 4**Isolation of the enzyme**

5 In order to isolate the enzyme, the culture medium of *Trichoderma reesei* (Rut C-30, VTT D-86271) was first treated with bentonite, as described by Zurbriggen et al. (1990). Then the solution was concentrated by ultrafiltration and the concentrate was dried by spray drying.

10 The isolation of the enzyme was started by dissolving the spray dried culture medium in a phosphate buffer. The insoluble material was separated by centrifugation and the enzyme solution was buffered by gel filtration to pH 7.2 (Sephadex G-25). The enzyme solution was pumped at this pH through a cation exchange chromatography column (CM-Sepharose FF), to which a part of the proteins of the sample were bound. The desired enzyme was collected in the fractions eluted through the column.

15 At said pH (pH 7.2) the enzyme solution was pumped to an anion exchange chromatography column (DEAE-Sepharose FF), to which most of the proteins of the sample were bound. The desired enzyme was collected in the fraction eluted through the column.

20 The enzyme-containing fractions were further purified by using hydrofobic interaction chromatography (Phenyl Sepharose FF). The enzyme was bound to said material at a salt concentration of 0.3 M $(\text{NH}_4)_2\text{SO}_4$. The bound enzyme was eluted with a buffer at pH 6.5, so as to form a decreasing linear concentration gradient of $(\text{NH}_4)_2\text{SO}_4$ from 0.3 to 0 M. After this, elution was continued with the buffer of pH 6.5. The mannanase enzyme was collected at the end of the gradient and in the fractions collected after that.

30 The enzyme solution was buffered by gel filtration to pH 4.3 (Sephadex G-25). The enzyme was bound at this pH to a cation exchange chromatography column (CM-Sepharose FF), and a part of the proteins bound to the column (i.a. most of the remaining cellulases) were eluted with a buffer, pH 4.4. The mannanase enzyme was eluted with a buffer, pH 4.3, to which sodium chloride was added in order to form a

linear concentration gradient of sodium chloride from 0 to 0.05 M. The purified enzyme was collected in the fractions eluted by the gradient.

5 **Example 5**
 Enzyme characterization

10 The protein properties of the enzyme preparation purified according to Example 1 were determined by methods known per se in protein chemistry. The molecular weights were determined by the SDS-PAGE-method, the accuracy of which is about ± 10 %.

 The preparation contains two mannanase isoenzymes, which biochemically and functionally proved to be almost identical. The isoenzymes are named EM 3 and EM 4. Their properties are described in Table 1.

Table 1. Properties of the *Trichoderma reesei* mannanase

Property	Unit	EM 3	EM 4
Molecular weight under denaturing conditions	kDa	51	53
Isoelectric point		4.5	5.4
Relative specific activity	percentage of the activity of the reference substance		
- reference ^a		100	100
- guar mannan		45	40
- Insoluble mannan of Corozo nut		7	9
- Soluble mannan of the Corozo nut		126	145
Optimum pH		3 - 5.3	3 - 5.3
Optimum temperature at for activity testing	°C	70	70
Hydrolysis products of β -1,4-mannan		M ₂ , M ₃	M ₂ , M ₃

^a: Locust bean gum

M₂: mannobiose

M₃: mannotriose

The N-terminal amino acid sequences of enzymes EM3 and EM4 have also been determined and they are indicated in sequence listings SEQ ID NO. 6 and SEQ ID NO, respectively.

Example 6

Mannanases secreted by *Trichoderma reesei* into its culture medium

The fungus *Trichoderma reesei* (VTT-D-86271) was cultivated in a fermentor for 5 days on a culture medium, whose main components were cellulose and corn steep

symp. After the cultivation the cells were separated from the solution by centrifugation and the mannanase activity of the culture medium was characterized by isoelectric focusing, ionexchange chromatography and chromatofocusing.

5 The isoelectric focusing was made by Phast equipment (Pharmacia-LKB) according to the instructions by the manufacturer on a gel supplied by the same manufacturer. A pH gradient from pH 3 to pH 9 was formed in the gel during focusing. After the isoelectric focusing time the proteins were transferred from this gel to a staining gel which contained Locust bean gum mannan. By staining with Congo red, the location of the
10 the mannanase proteins could be detected on this gel, which made it possible to estimate the isoelectric point of the mannanase enzymes.

Each isoenzyme's relative portion of the total mannanase activity isoenzyme was estimated by running chromatofocusing on a PBE 94 gel supplied by Pharmacia-LKB
15 according to the instructions of the manufacturer. The mannanase activity of the fractions obtained was measured and they were then subjected to an activity staining as described above. Furthermore the different fractions of the purification procedure were, if needed, subjected to activity staining in addition to mannanase determination. These methods made it possible to make an estimation of the relative amounts of the enzymes
20 in the solution of the Example with an accuracy of 10 to 20 %.

Table 2 indicates the isoelectric points of the mannanase enzymes and their relative activities.

Table 2. Isoelectric points of the *Trichoderma reesei* mannanases and their relative portions of the total mannanase activity

	pI	Estimated portion of the mannanase activity (%)
EM 1	3.8	5 - 15
EM 2	4.1	10 - 30
EM 3	4.5	30 - 40
EM 4	5.4	30 - 40
EM 5	6.5	< 5

Example 7

Hydrolysis of cellulosic pulp

Pine kraft pulp, pine saw dust and glucomannan fractions isolated from them were hydrolysed with *T. reesei* mannanase. The enzyme dosage was 5000 nkat/g calculated on basis of the glucomannan. The treatment was carried out in 50 mM citrate buffer at pH 5.0. The treatment lasted for 24 h and the temperature was 45 °C. After the treatment the dissolved sugars were determined after acid hydrolysis. The result is indicated in percent of the original glucomannan which was hydrolyzed. (Table 3).

Table 3. The relative amounts of hydrolyzed glucomannans

Substrate	percentage of hydrolyzed glucomannan
pine saw dust	11
mannan isolated from pine tree	65
pine kraft pulp	10
mannan isolated from pine kraft pulp	79

10

Thus, the mannanase are efficient in hydrolyzing mannan isolated from both pine tree and pine kraft pulp. As a result of the hydrolysis, glucomanno-oligomers are formed.

15

Example 8**Bleaching of cellulose pulp (mannanase pretreatment)**

20

Pine kraft pulp (kappa 26.2) was treated with *T. reesei* mannanase at 5 % consistency for 2 h at pH 5 and 45 °C. After the treatment the pulp was washed (2x10xdry matter of the pulp) and bleached with a 1-stage peroxide sequence. Subsequent to the peroxide stage the pulp was acidified and its brightness, kappa and viscosity were determined. The results are indicated in Table 4. The chemical dosages and conditions were: H₂O₂ 3 %, NaOH 1.5 %, DTPA 0.2 %, MgSO₄ 0.5 %, 80 °C, 1 h.

25

Table 4. Peroxide bleaching of kraft pulp subsequent to mannanase treatment

Dose nkat/g	Released reducing sugars in % of dry matter	Kappa	Bright- ness, %	Viscosity dm ³ /kg
100	0.25	13.9	42.9	970
500	0.48	13.5	43.2	940
ref	0	17.6	41.0	980

After a simple mannanase treatment, the kappa reduction during the peroxide stage was 23 % better than in the reference test.

Example 9

Bleaching of cellulose pulp

The procedure described in Example 5 was followed with the difference that coniferous pulp was treated with mannanase and xylanase. The pulp was subjected to peroxide bleaching similar to the one of Example 4. The results appear from Table 5.

Table 5. Bleaching with peroxide of kraft pulp after mannanase/xylanase treatment

Enz.	Dose nkat/g	Reducing sugars in % of dry matter	Kappa	Bright- ness, %	Viscosity dm ³ /kg
XYL + MAN	100 + 100	0.45	12.8	45.0	990
XYL + MAN	100 + 500	0.9	11.8	44.8	950
XYL	100	0.41	16.8	43.7	990
ref	-	0	17.6	41.0	980

XYL = xylanase

MAN = mannanase

Combination of the mannanase treatment with a xylanase treatment lowered the kappa of the peroxide bleached pulp by 32 % compared to the reference treatment.

Example 10

Bleaching test carried out for cellulose pulp

Pine kraft pulp cooked in a laboratory (kappa 34.4) was treated with mannanase purified from *T. reesei* at a consistency of 5 %, at pH 5 and at a temperature of 45 °C. After the treatment the pulp was washed and bleached with the sequence (D50/C50)EDED.

Table 6. Bleaching with chlorine chemicals of coniferous pulp treated with enzymes

Enz.	Dose nkat/g	Chlori- nation factor	Int. kappa	Bright- ness, %	Viscosity dm ³ /kg
MAN	90	0.15	6.4	88.4	1050
XYL pI9	100	0.15	5.6	89.5	1080
MAN + XYL	90 + 100	0.15	5.2	90.3	1060
ref	-	0.15	6.5	88.1	1050
ref	-	0.18	4.5	90.5	1050

In comparison to a simple xylanase treatment, a combined xylanase and mannanase treatment provided a considerably much smaller intermittent kappa and a higher final brightness.

Example 11

Bleaching of mannanase-treated sulphite pulp

Coniferous sulphite pulp was treated with mannanase at 5 % consistency at pH 5 and a temperature of 45 °C for 2 h. After the treatment the pulp was washed and bleached with a one-stage peroxide sequence. After the peroxide stage the brightness, kappa and viscosity of the pulp were determined. The chemical dosages were: 1.5 % H₂O₂, 1.2 % NaOH 1,2 % and 0.05 % MgSO₄. The bleaching time was 60 min at 80 °C. (Enz = XYL 100, XYL 100+500, MAN 500)

It was found that the mannanase treatment made it possible to improve the bleachability of sulphite pulp.

Example 12**Bleaching test of kraft pulp treated with mannanase and xylanase**

5 Coniferous kraft pulp (spruce/pine) was treated with mannnase and xylanase as in Example 7. After the treatment the pulp was washed and bleached by the sequence D(ED)DED. The bleaching conditions are shown in Table 7A.

10 After the mannanase treatment the intermittent kappa and the brightness were better than in the reference pulp. When the mannanase treatment was combined with a xylanase treatment the intermediate kappa was much lower and an over 90 % final brigthness was obtained with a 15 % saving in chemical consumption (Table 7B).

Table 7A. Bleaching conditions. Bleaching sequence: D(EP)DED). Chlorination factors 0.15 and 0.20.

Stage	Consistency %	T °C	Duration min.	Final pH	Act. Cl %	P	NaOH
D0	3	60	60	2.0 - 1.9	3.3 ^{1,3)} 4.3 ²⁾	%	%
EP	10	80	60	10.4 - 10.5		0.3	1.7 ^{1,3)} 2.1 ²⁾
D1	10	70	180	3.4 - 3.6	2.5 ^{1,2)} 3.0 ³⁾		
E2	10	70	60	10.8 - 10.9			0.9
D2	10	70	180	3.8 - 4.1			0.5 - 1.0 ^{1,3)} 0.7

1) Enzyme treated pulp, chlorination factor 0.15

2) Chlorination factor 0.20

3) Reference, chlorination factor 0.15

Table 7B. Chlorine bleaching of mannanase and xylanase treated pulps. SequenceD(EP)DED. Initial kappa of pulp: 21.6.

Enzyme	Dosage, nkat/g	Reducing sugars, % of dry matter	Chlorination factor	Intermediate kappa	D-1 brightness %	Final brightness %	Viscosity dm ³ /kg	Chemicals consumption % act. Cl
MAN	500	0.40	0.15	6.2	80.1	89.8	940	6.7
XYL	100	0.38	0.15	5.5	82.5	90.2	980	6.4
MAN + XYL	500 + 100	0.70	0.15	5.2	83.5	90.7	960	6.3
ref	-	-	0.15	6.5	79.8	89.7	970	7.1
ref	-	-	0.20	5.1	82.8	90.1	970	7.4

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SEQUENCE LISTING

(1) GENERAL INFORMATION:

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(ii) TITLE OF INVENTION: Mannanase enzymes, genes coding for them and a method for isolating the genes, as well as a process for bleaching of lignocellulosic pulp

(iii) NUMBER OF SEQUENCES: 7

(iv) PRIOR APPLICATION DATA:

- (A) APPLICATION NUMBER: FI 922373
- (B) FILING DATE: 22 MAY 1992

- (A) APPLICATION NUMBER: FI 931193
- (B) FILING DATE: 17 MARCH 1993

(2) INFORMATION FOR SEQ ID NO: 1:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 536 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: Trichoderma reesei
- (B) STRAIN: QM9414

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:

GAATTCGGCA CGAGAGTCTC TCCTTTTTTT CTCACTATAT CTTTCCTTTG TCCATGTTGC	60
TGTTGTCGTC GTTGTTCCTG AGACTATGCT GGCCTGTTG GTCCGGATGA ACTGGTGTT	120
GGGATGGCTG GAGCCTGTGG AAGCGGGCTC GCCGTGGCGT TGGTCAACCG ATTGTATATC	180
AGTCTATGCC TCTGTACACT TCGTCTCTAG CGAAGAGGAG TTGAATACAA ATCTGTAAAA	240
CACCTGACTG TGTCTTCTAG CTATGAGACT CCCTTGCCTA CTGGAGCCTT CAAGATACTT	300
TGGTACTGTA TGAGACCACG CCTACCTCGA CTCATGTTT GAAACCAGTC AGTAATTCTC	360
TATGAACATG AAACAACACA TTGATCTCTG TAACATCTCA TTGCATAGTA AACCTTCTTA	420
CATTGATTAC TGGCTATGAA CAAAGGTTGT AGGGTAGGTA ACGAAAAAAAA AAAAAAAAAA	480
AAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA CTCGAG	536

(2) INFORMATION FOR SEQ ID NO: 2:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 243 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: *Trichoderma reesei*
- (B) STRAIN: QM9414

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2:

GAATTCGGCA CGAGCTCTTA ACCAACCACC AACTACAGC CACCCACCAT GTCCGCCCAA	60
GACTACTACG TCCAGCGCCT CCGGAGGCAG CAGCAACGGT TATCCTCCTC AGCAGTACCA	120
CCAGCAGCAG CAACAGCCAT ACGGCCAGCA GCAGCCAGGG TATGACCAGC AGTACCCCGG	180
CGCCGGAGTC GATGGAGGCC CCGACGGAGA GCGCGGCCTC GCGCGACCC TCGTCGGCGG	240
CGG	243

(2) INFORMATION FOR SEQ ID NO: 3:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 289 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: *Trichoderma reesei*
- (B) STRAIN: QM9414

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3:

```
CTGCAGCCGC CGGAGCCATT GGGGCGAATC TCATTGAGAA CGCATACAAG AAGCACAAGG      60
AGGAAAAGAT GTACAATGAT GGC GGACACC ATGGCCATCA CCACCATTGG CATCATCATA      120
AGCACTAGAA TGATTTGATG AAGAATATGA CATTGCATGC CTGCTACATA CGTAGATTAT      180
GATTGGGGGA GGGGCGTTTT TGATGAATAT ATATATATAT CAATAAAAAA AAAAAAAAAA      240
AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAC TCGAG                      289
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(2) INFORMATION FOR SEQ ID NO: 4:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 865 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: *Trichoderma reesei*
- (B) STRAIN: QM9414

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 4:

GAATTCGGCA CGAGTTTTTT TTTTTTTTTT TACAGTATAC GTTAGGTGTA TTCGAACGAA	60
AGCCTGAAAT AGGTACAAAT AGCCACAACA GAAACGATTG TCTGACCTCC ACTGAGAAAA	120
TCCCCCTCGT CCCCCAGACA GGTCTGATCA AACACCAGCT ATCTACCTGC AAATGCCCCA	180
TATGTAGGTC ACACAGTAAG GAGTGGATTC GCTATATACA TCGGGTACGT TCGCATCACC	240
CCATGGCAAG GGAGGTGACT TAAGCAAAAC CGCCACTAAC CACAAAGCTC AACTGCATAG	300
TATCGACTTC AAGGAAAACA CGGACAAATA ATCATCATGG TTGCCTTTTG CAGCCTCATC	360
TGCGCTCTCA CGAGGCATCG CCAGTACTCT GGCGATGCCC ACAGGCTCGA GCCTGAGAGC	420
AGTGTCACG TCACAGAGCG TGGCATGTAC GACTTTGTTC TTGGAGCTCA CAATGATCAT	480
CGCCGTCGTG CTAGCATCAA CTACGACCAA AACTACCAA CTGGCGGACA AGTCAGCTAT	540
TGCGCTTCCA AACTGGCTT CTCAGTGAAC TGGAACACTC AAGATGACTT TGTTGTGGGC	600
GTTGGTTGGA CGACTGGATC TTCTGCTCCC ATCAACTTTG GCGGCTCTTT TAGTGTCAAC	660
AGCGGAACTG GCCTGCTTTC CGTCTATGGC TGGAGCACCA ACCCACTGGT TGAGTACTAC	720
ATCATGGAGG ACAACCACAA CTACCCAGCA CAGGGCACCG TCAAGGGAAC CGTCAACAGC	780
GACGGGGCAC TTACACCATC TTGGGGGATT ACCGTGTAAC GAGCCTTCCA TCCAGGGTAC	840
AGCGCCTTCA ACCGTACATT TCCGT	865

(2) INFORMATION FOR SEQ ID NO: 5:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 319 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: *Trichoderma reesei*
- (B) STRAIN: QM9414

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:

CAGATGAACT ACCCAGGTTG TCGCTGTCGA AAGGCTGGGG TGGTAGTGGT TCTGCCTCAC	60
AGAGTGTGAG CAACTAGGTT CTGTTGATGT TGAAGTGGAG TGGATGAGGG GTTTGAGCTG	120
GTATGTAGTA TTGGGGTGGT TAGTGAGTTA ACTTGACAGA CTGCACTTTG GCAACAGAGC	180
CGACGATTAA GAGATTGCTG TCATGTAAGT AAAGTAGCCT GCCTTTGACG CTGTATGCTC	240
ATGATACATG CGTGACATCG AAATATATCA GCCAAAGTAT CCGTCCGGCG AAAAAAAAAA	300
AAAAAAAAAA AACTCGAG	319

(2) INFORMATION FOR SEQ ID NO: 6:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 15 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(iv) FRAGMENT TYPE: N-terminal

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 6:

Xaa	Xaa	Xaa	Phe	Val	Thr	Ile	Ser	Xaa	Thr	Gln	Xaa	Xaa	Ile	Asp
1				5					10					15

(2) INFORMATION FOR SEQ ID NO: 7:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 15 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(iv) FRAGMENT TYPE: N-terminal

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 7:

Xaa	Xaa	Xaa	Xaa	Xaa	Thr	Ile	Ser	Gly	Thr	Gln	Xaa	Xaa	Ile	Asp
1					5					10				15

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>11</u> , line <u>30</u>	
B. IDENTIFICATION OF DEPOSIT	
Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution DSM-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH	
Address of depositary institution (including postal code and country) Mascheroder Weg 1 B, D-3300 Braunschweig, Federal Republic of Germany	
Date of deposit 30 December 1992	Accession Number DSM 7363
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
In respect of those designations in which a European patent is sought, a sample of the deposited microorganism will be made available only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28(4) EPC) until the publication of the mention of the grant of the European patent or until the date on which the application has been refused or is deemed to be withdrawn.	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>11</u> , line <u>30</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution DSM-DEUTSCHE SAMMLUNG VON MIKROORGANISMEN UND ZELLKULTUREN GmbH	
Address of depositary institution (including postal code and country) Mascheroder Weg 1 B, D-3300 Braunschweig, Federal Republic of Germany	
Date of deposit 30 December 1992	Accession Number DSM 7364
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
In respect of those designations in which a European patent is sought, a sample of the deposited microorganism will be made available only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28(4) EPC) until the publication of the mention of the grant of the European patent or until the date on which the application has been refused or is deemed to be withdrawn.	
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

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Address of depositary institution (including postal code and country) Mascheroder Weg 1 B, D-3300 Braunschweig, Federal Republic of Germany	
Date of deposit 30 December 1992	Accession Number DSM 7366
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D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

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A. The indications made below relate to the microorganism referred to in the description on page <u>11</u> , line <u>30</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
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Address of depositary institution (including postal code and country) Mascheroder Weg 1 B, D-3300 Braunschweig, Federal Republic of Germany	
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In respect of those designations in which a European patent is sought, a sample of the deposited microorganism will be made available only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28(4) EPC) until the publication of the mention of the grant of the European patent or until the date on which the application has been refused or is deemed to be withdrawn.	
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Claims:

1. A DNA sequence, c h a r a c t e r i z e d in that it codes for a *Trichoderma* enzyme having mannanase activity and transferred into a yeast or fungus strain induces that strain to produce mannanase.
5
2. The DNA sequence according to claim 1, wherein the DNA sequence hybridizes to the DNA sequence of SEQ ID NO. 1.
- 10 3. A vector, c h a r a c t e r i z e d in that it contains a DNA sequence of claim 1 or claim 2.
4. A yeast vector, c h a r a c t e r i z e d in that it is selected from the group comprising pMAN1, pMAN2, pMAN3 and pMAN4.
15
5. A recombinant yeast or fungus strain, c h a r a c t e r i z e d in that it contains a DNA sequence according to claim 1 or claim 2 and that it codes for mannanase.
6. The yeast strain according to claim 5, c h a r a c t e r i z e d in that it is
20 *Saccharomyces cerevisiae*.
7. The yeast strain according to claim 6, c h a r a c t e r i z e d in that it is selected from the group comprising S.c.man1, S.c.man2, S.c.man3 and S.c.man4, and it is deposited under one of the deposition numbers DSM 7363, DSM 7364, DSM 7366
25 wherein DSM 7365, respectively, in the despository Deutsche Sammlung von Mikroorganismen und Zellkulturen GmbH.
8. The yeast strain according to claim 5, c h a r a c t e r i z e d in that it belongs to the species *Trichoderma*.
30
9. A method for isolating a genes coding for mannanases, c h a r a c t e r i z e d by
- enriching the messenger RNA pool of a microorganism producing mannanase

activity in respect of the messenger RNA by culturing the microorganism in conditions which will induce the mannanase production of said microorganism,

- isolating messenger RNA from the microorganism strain and preparing cDNA corresponding to the isolated mRNA,
- 5 - placing the cDNA thus obtained in a vector under the control of a yeast promoter,
- transforming the recombinant plasmids into a yeast strain which naturally does not produce the corresponding mannanase in order to provide an expression library,
- cultivating the yeast clones thus obtained on a culture medium in order to express the expression library in the yeast,
- 10 - separating the yeast clones producing the mannanase from the other yeast clones,
- isolating the plasmid-DNA of said yeast clones, and,
- if desired, sequencing the DNA in order to determine the DNA sequence corresponding to the mannanase.

15 10. The method according to claim 9, wherein the microorganism is a fungus.

11. The method according to claims 9 or 10, wherein gene library is constructed from a fungus strain belonging to the genus *Trichoderma*.

20 12. The method according to claim 11, wherein the gene library is constructed from a fungus of the species *Trichoderma reesei*.

25 13. The method according to claim 10, wherein the fungus is cultivated on a medium containing as carbon source at least mannanase- and/or cellulose-containing substrate and possibly acetylglucuronoxylan and/or xylan.

14. The method according to claim 9, wherein the yeast strain used belongs to the genus *Saccharomyces*.

30 15. The method according to claim 14, wherein the yeast strain comprises a *Saccharomyces cerevisiae* strain which is cultivated on a glucose-containing culture medium in order to produce mannanase.

16. The method according to any one of claims 9 to 15, wherein the cDNA produced from the messenger RNA of the microorganism is ligated to a plasmid vector, in which the yeast promoter comprises PGK.

5 17. The method according to claim 16, wherein the plasmid vector comprises the vector pAJ401.

18. The method according to any one of claims 9 to 17, wherein the recombinant yeast clones are detected on basis of their mannanase activity by enzyme activity test carried
10 out in solution or by plate tests.

19. A method for constructing yeast strains capable of expressing mannanase,
c h a r a c t e r i z e d b y

- isolating the DNA sequences coding for mannanase from a suitable fungus strain,
- 15 - forming a yeast vector from the DNA sequence, and
- transferring the yeast vector to a suitable fungus strain.

20. The method according to claim 23, wherein the yeast strains are transferred to yeast strain of the genera *Saccharomyces*, *Kluyveromyces*, *Pichia* or *Hansenula*.

20

21. The method according to claim 20, wherein the yeast vectors are transferred to a *Saccharomyces cerevisiae* strain.

22. A process for producing mannanase enzyme, c h a r a c t e r i z e d b y

- 25 - isolating from a *Trichoderma reesei* fungus a DNA sequence coding for mannanase,
- forming a vector containing said DNA sequence,
- transferring the obtained vector to a suitable yeast or fungus strain in order to provide a recombinant stain,
- 30 - cultivating the recombinant strain under conditions in which the strain is capable of expressing mannanase, and
- recovering the produced mannanase.

23. A process for producing a mannanase enzyme or an enzyme mixture,
c h a r a c t e r i z e d by

- treating the culture medium of a fungus strain belonging to the species
Trichoderma reesei with bentonite,
- 5 - subjecting the treated culture medium first to cation exchange and then to anion
exchange essentially at pH 7 to 7.5,
- collecting the elutes of the ionexchange and purifying them on a chromatography
column with hydrophobic interaction chromatography, and
- 10 - optionally subjecting the enzyme product to additional treatment for purification
thereof.

24. An enzyme preparation, c h a r a c t e r i z e d in that it contains at least one
mannanase produced by a fungus of the genus *Trichoderma* and selected from the group
comprising an enzyme having mannanase activity and an isoelectric point (pI) of about
15 3.8, an enzyme having mannanase activity and a pI of about 4.1, an enzyme having
mannanase activity and a pI of about 4.5, an enzyme having mannanase activity and a
pI of about 5.4, and an enzyme having mannanase activity and a pI of about 6.5, the
isoelectric points being determined by isoelectric focusing.

20 25. A process for hydrolyzing mannopolymers, c h a r a c t e r i z e d by using in
the hydrolysis the mannase of a fungus of the genus *Trichoderma*.

26. The process according to claim 25, wherein an enzyme preparation containing at
least one mannanase is used in the hydrolysis, said preparation being produced by a
25 fungus of the species *Trichoderma reesei*, by a strain derived thereof, or by another
production host to which the genes coding for the production of mannanases have been
transferred.

27. The process according to claim 26, wherein the enzyme preparation is produced by
30 a yeast strain of the species *Saccharomyces cerevisiae* to which a gene coding for
mannanase production has been transferred.

28. The process according to claim 26, wherein glucomannans of cellulose pulp are hydrolyzed.
- 5 29. The process according to claim 28, wherein the mannanase is used together with at least one xylan.
30. The process according to claim 28, wherein glucomannans of cellulose pulp are hydrolyzed.
- 10 31. A process for bleaching of cellulose pulp, characterized by treating the pulp in connection with the bleaching with a *Trichoderma* mannanase.
32. The process according to claim 31, wherein the mannanase treatment is carried out before the bleaching.
- 15 33. The process according to claim 32, wherein the mannanase treatment is carried out before a chlorine free bleaching of pulp.
34. The process according to claim 33, wherein the mannanase treatment is carried out before peroxide bleaching.
- 20 35. The process according to claim 32, wherein the mannanase treatment is combined with xylanase treatment.
- 25 36. The process according to any one of claim 31 to 35, wherein the mannanase treatment is carried out in connection with bleaching of sulphite pulp.

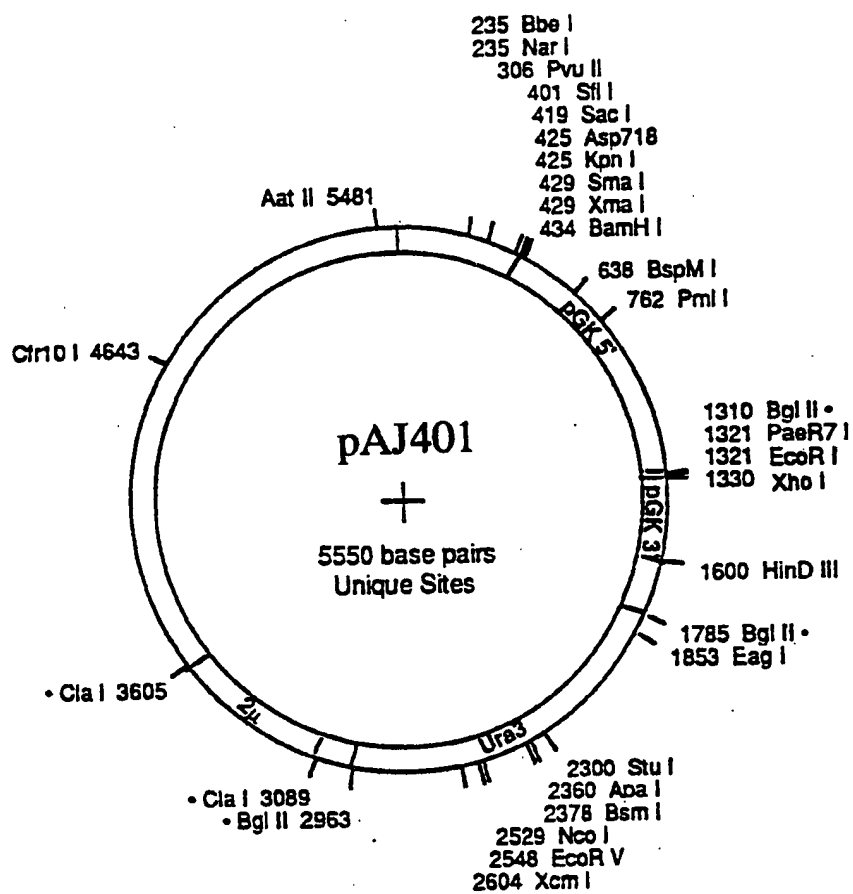


FIG. 1

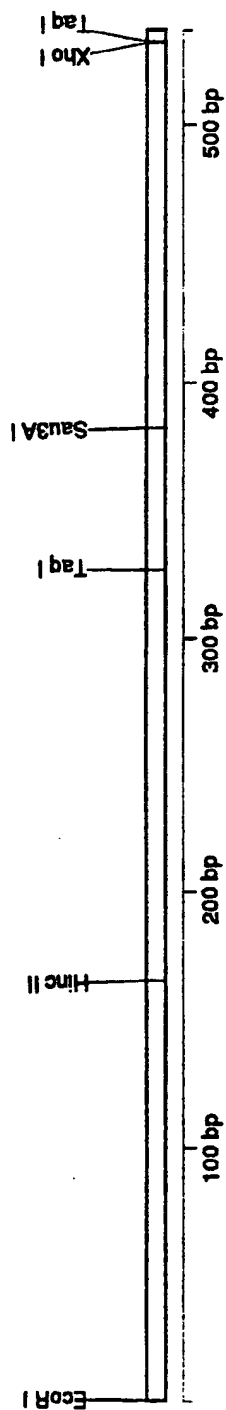


FIG. 2

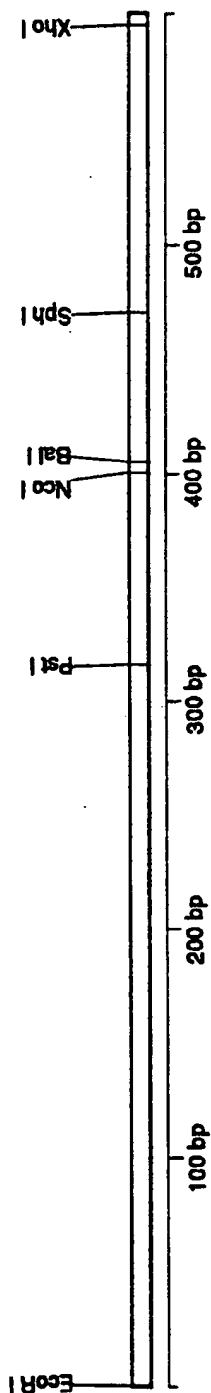


FIG. 3

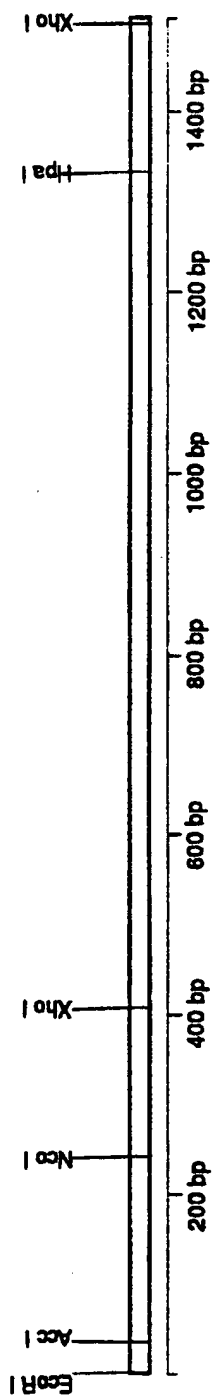


FIG. 4

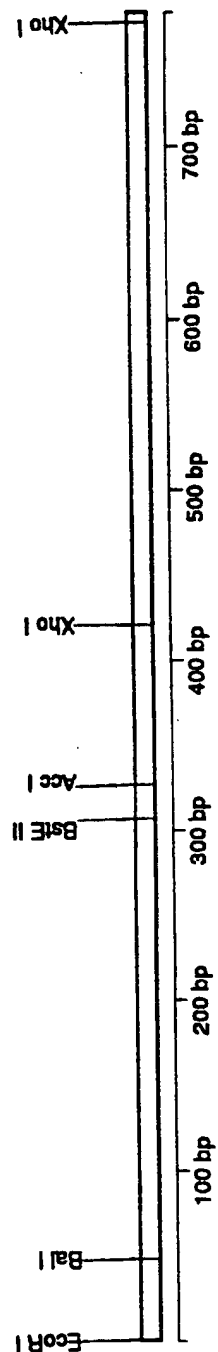


FIG. 5

INTERNATIONAL SEARCH REPORT

International application No.

PCT/FI 93/00219

A. CLASSIFICATION OF SUBJECT MATTER		
IPC5: C12N 9/42, C12N 15/56 // C12N 9/42, C12R 1:885 According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED		
Minimum documentation searched (classification system followed by classification symbols)		
IPC5: C12N		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
SE,DK,FI,NO classes as above		
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	Dialog Information Services, File 357, Biotechnology Abstracts, Dialog accession no. 146065, DBA accession no. 93-04117, Torrie J P et al: "The substrate specificity of an exocellular beta mannanase from <i>Trichoderma harzianum</i> E58 - endo-1,4-beta-D-mannanase purification and characterization for potential application in pulping (conference paper), Biodeterior Biodegr. 8, 542-44 1991 --	1-36
X	Dialog Information Services, File 5, BIOSIS, Dialog accession no. 8150876, BIOSIS Number. 91071876, Torrie J P et al: "Production of beta mannanases by <i>trichoderma harzianum</i> E58", Appl microbiol biotechnol 34 (3) 1990, 303-307 --	1-36
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. <input checked="" type="checkbox"/> See patent family annex.		
<p>* Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&" document member of the same patent family</p>		
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31 August 1993		08 -09- 1993
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International application No.

PCT/FI 93/00219

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO, A1, 9118974 (CHEMGEN CORPORATION), 12 December 1991 (12.12.91) -----	1-36

30/07/93

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